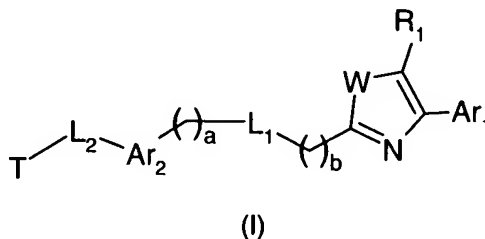


Amendments to the Claims

Please amend the claims as follows. This listing of claims will replace all prior versions and listings of claims in the application.

1. (Currently Amended) A compound of Formula (I):



wherein

a and b are, independently, equal to 0, 1, or 2, wherein the values of 0, 1, and 2 represent a direct bond, $-\text{CH}_2-$, and $-\text{CH}_2\text{CH}_2-$, respectively, and wherein the $-\text{CH}_2-$ and $-\text{CH}_2\text{CH}_2-$ groups are optionally substituted 1 to 2 times with a substituent group, wherein said substituent group(s) are selected from the group consisting of: -alkyl, -aryl, -alkylene-aryl, -arylene-alkyl, -alkylene-arylene-alkyl, -O-alkyl, -O-aryl, and -hydroxyl;

W is $-\text{O}-$, $-\text{S}-$, or $-\text{N}(\text{R}_2)-$,

wherein

R_2 is

- a) -hydrogen;
- b) -alkyl;
- c) $-\text{L}_3\text{-D-G}$
- d) $-\text{L}_3\text{-D-alkyl}$;
- e) $-\text{L}_3\text{-D-aryl}$;
- f) $-\text{L}_3\text{-D-heteroaryl}$;
- g) $-\text{L}_3\text{-D-cycloalkyl}$;
- h) $-\text{L}_3\text{-D-heterocyclyl}$;
- i) $-\text{L}_3\text{-D-arylene-alkyl}$;
- j) $-\text{L}_3\text{-D-alkylene-arylene-alkyl}$;

- k) – L₃-D-alkylene-aryl;
- l) – L₃-D-alkyl-G;
- m) – L₃-D-aryl-G;
- n) – L₃-D-heteroaryl-G;
- o) – L₃-D-cycloalkyl-G;
- p) – L₃-D-heterocyclyl-G;
- q) – L₃-D-arylene-alkyl-G;
- r) – L₃-D-alkylene-arylene-alkyl-G; or
- s) – L₃-D-alkylene-aryl-G;

wherein

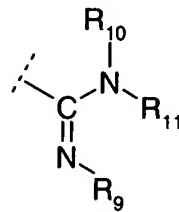
L₃ is a direct bond, –alkylene, –alkenylene, or alkynylene;

D is a direct bond, –CH₂–, –O–, –N(R₅)–, –C(O)–, –CON(R₅)–, –N(R₆)C(O)–, –N(R₆)CON(R₅)–, –N(R₅)C(O)O–, –OC(O)N(R₅)–, –N(R₅)SO₂–, –SO₂N(R₅)–, –C(O)–O–, –O–C(O)–, –S–, –S(O)–, –S(O₂)–, or –N(R₅)SO₂N(R₆)–, –N=N–, or –N(R₅)–N(R₆)–;

wherein

R₅ and R₆ are independently selected from the group consisting of: –hydrogen, –alkyl, –aryl, –arylene-alkyl, –alkylene-aryl, and –alkylene-arylene-alkyl; and

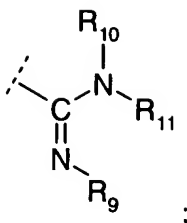
G is hydrogen, –CN, –SO₃H, –P(O)(OH)₂, –P(O)(O-alkyl)(OH), –CO₂H,



–CO₂-alkyl, an acid isostere, –NR₇R₈, or

wherein

R₇ and R₈ are independently selected from the group consisting of: –hydrogen, –alkyl, –L₄-E-alkyl, –L₄-E-aryl, –C(O)-alkyl, –C(O)-aryl, –SO₂-alkyl, –SO₂-aryl, and



wherein

R_9 , R_{10} , and R_{11} are independently selected from the group consisting of: -hydrogen, -alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, and -alkylene-arylene-alkyl;

L_4 is a direct bond, -alkylene, -alkenylene, or -alkynylene;

E is a direct bond, - CH_2 -, -O-, - $N(R_{12})$ -, -C(O)-, -CON(R_{12})-, - $N(R_{12})C(O)$ -, - $N(R_{12})CON(R_{13})$ -, - $N(R_{12})C(O)O$ -, -OC(O) $N(R_{12})$ -, - $N(R_{12})SO_2$ -, - $SO_2N(R_{12})$ -, -C(O)-O-, -O-C(O)-, -S-, -S(O)-, -S(O)₂-, - $N(R_{12})SO_2N(R_{13})$ -, -N=N-, or - $N(R_{12})N(R_{13})$ -

wherein

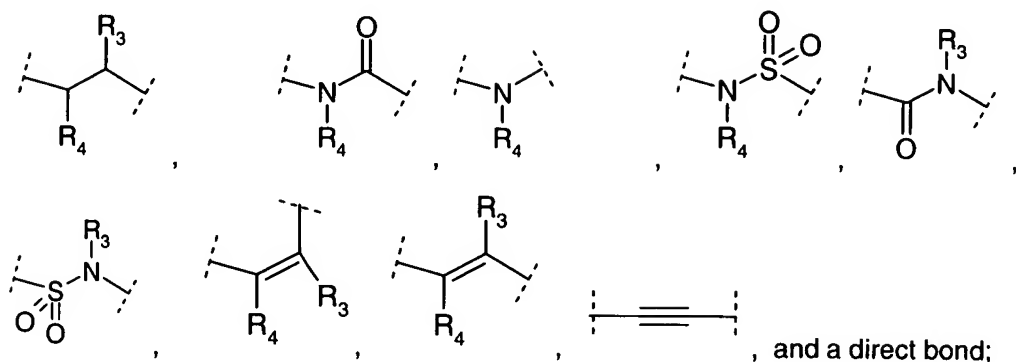
R_{12} and R_{13} are independently selected from the group consisting of: -hydrogen, -alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, and -alkylene-arylene-alkyl;

R_1 is

- a) -hydrogen;
- b) -fluoro;
- c) -chloro;
- d) -bromo;
- e) -iodo;
- f) -cyano;
- g) -alkyl;
- h) -aryl;
- i) -alkylene-aryl;
- j) -heteroaryl;
- k) -alkylkene-heteroaryl;

- l) -cycloalkyl;
- m) -alkylene-cycloalkyl
- n) - heterocyclyl; or
- o) - alkylene-heterocyclyl;

L₁ is selected from the group consisting of:



wherein R₃ and R₄ are independently selected from the group consisting of: hydrogen, chloro, fluoro, bromo, alkyl, aryl, -alkylene-aryl, -cycloalkyl, -alkylene-cycloalkyl, -heterocyclyl, -alkylene-heterocyclyl, and -alkynylene [.] ;

Ar₁ is an aryl, heteroaryl, fused cycloalkylaryl, fused cycloalkylheteroaryl, fused heterocyclylaryl, or fused heterocyclylheteroaryl group optionally substituted 1 to 7 times;

Ar₂ is an arylene, heteroarylene, fused arylcycloalkylene, fused cycloalkylarylene, fused cycloalkylheteroarylene, fused heterocyclylarylene, or fused heterocyclylheteroarylene group optionally substituted 1 to 7 times;

L₂ is selected from the group consisting of: -CH₂-, -O-, alkylene, alkenylene, alkynylene, -K-alkylene-, -alkylene-K-, -alkylene-K-alkylene-, -alkenylene-K-alkylene-, -alkylene-K-alkenylene-, -arylene-K-alkylene-, alkylene-K-arylene, -heteroarylene-K-alkylene-, alkylene-K-heteroarylene, -arylene-K-, -K-arylene-, -heteroarylene-K-, -K-heteroarylene, and a direct bond,

wherein

K is a direct bond, $-N(R_{20})-$, $-C(O)-$, $-CON(R_{20})-$, $-N(R_{20})C(O)-$,
 $-N(R_{20})CON(R_{21})-$, $-N(R_{20})C(O)O-$, $-OC(O)N(R_{20})-$, $-N(R_{20})SO_2-$, $-$
 $SO_2N(R_{20})-$, $-C(O)-O-$, $-O-C(O)-$, $-S-$, $-S(O)-$, $-S(O_2)-$,
 $-N(R_{20})SO_2N(R_{21})-$, $-N=N-$, or $-N(R_{20})-N(R_{21})-$; $-N(R_{20})-$, $-C(O)-$, $-$
 $CON(R_{20})-$, $-N(R_{20})C(O)-$, $-N(R_{20})CON(R_{21})-$, $-N(R_{20})C(O)O-$,
 $-OC(O)N(R_{20})-$, $-N(R_{20})SO_2-$, $-SO_2N(R_{20})-$, $-C(O)-O-$, $-O-C(O)-$, $-S$, $-$
 $S(O)-$, $-S(O_2)-$, $-N(R_{20})SO_2N(R_{21})-$, $-N=N-$, or $-N(R_{20})-N(R_{21})-$ or a
direct bond,

wherein

R_{20} and R_{21} are independently selected from the group: -hydrogen,
-alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, and -alkylene-
arylene-alkyl;

T is selected from the group consisting of: hydrogen, alkyl, cycloalkyl, heterocyclyl,
aryl, heteroaryl, fused cycloalkylaryl, fused cycloalkylheteroaryl, fused
heterocyclylaryl, and fused heterocyclylheteroaryl group optionally substituted 1 to 7
times,

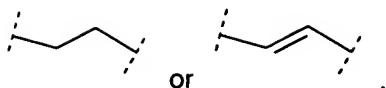
or a pharmaceutically acceptable salt, solvate, or prodrug thereof.

2. (Currently Amended) The compound of Formula (I) according to claim
1 or a pharmaceutically acceptable salt thereof, wherein W is $-O-$ or $-N(R_2)-$, wherein
 R_2 is hydrogen, alkyl, or $-L_3$ -D-alkylene-aryl, wherein L_3 is alkylene, and D is
 $-CO(NR_5)-$, wherein R_5 is hydrogen.

3. (Currently Amended) The compound of Formula (I) according to claim
1 or a pharmaceutically acceptable salt thereof, wherein R_1 is hydrogen or aryl.

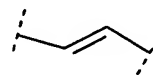
4. (Currently Amended) The compound of Formula (I) according to claim
1 or a pharmaceutically acceptable salt thereof, wherein R_1 is hydrogen.

5. (Currently Amended) The compound of Formula (I) according to claim
1 or a pharmaceutically acceptable salt thereof, wherein L_1 is



6. (Currently Amended) The compound of Formula (I) according to claim

1 or a pharmaceutically acceptable salt thereof, wherein L_1 is



7. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein Ar_1 is a phenyl or naphthyl group optionally having 1 to 5 substituents, wherein the substituents are independently selected from the group consisting of:

- a) -fluoro;
- b) -chloro;
- c) -bromo;
- d) -iodo;
- e) -cyano;
- f) -nitro;
- g) -perfluoroalkyl;
- h) - $J-R_{14}$;
- i) -alkyl;
- j) -aryl;
- k) -heteroaryl;
- l) -heterocyclyl;
- m) -cycloalkyl;
- n) - L_5 -aryl;
- o) - L_5 -arylene-aryl;
- p) - L_5 -arylene-alkyl;
- q) -arylene-alkyl;
- r) -arylene-arylene-alkyl;
- s) -J-alkyl;
- t) -J-aryl;
- u) -J-alkylene-aryl;
- v) -J-arylene-alkyl;

- w) -J-alkylene-arylene-aryl;
- x) -J-arylene-arylene-aryl;
- y) -J-alkylene-arylene-alkyl;
- z) -L₅-J-alkylene-aryl;
- aa) -arylene-J-alkyl;
- bb) -L₅-J-aryl;
- cc) -L₅-J-heteroaryl;
- dd) -L₅-J-cycloalkyl;
- ee) -L₅-J-heterocyclyl;
- ff) -L₅-J-arylene-alkyl;
- gg) -L₅-J-alkylene-arylene-alkyl;
- hh) -L₅-J-alkyl;
- ii) -L₅-J-R₁₄;
- jj) -arylene-J-R₁₄; and
- kk) -hydrogen;

wherein

L₅ is a direct bond, -alkylene, -alkenylene, or -alkynylene;

J is a direct bond, -CH₂-, -O-, -N(R₁₅)-, -C(O)-, -CON(R₁₅)-, -N(R₁₅)C(O)-, -N(R₁₅)CON(R₁₆)-, -N(R₁₅)C(O)O-, -OC(O)N(R₁₅)-, -N(R₁₅)SO₂-, -SO₂N(R₁₅)-, -C(O)-O-, -O-C(O)-, -S-, -S(O)-, -S(O₂)-, -N(R₁₅)SO₂N(R₁₆)-, -N=N-, or -N(R₁₅)-N(R₁₆)-,

wherein

R₁₄, R₁₅, and R₁₆ are independently selected from a group consisting of: -hydrogen, -alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, and -alkylene-arylene-alkyl.

8. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein Ar₁ is a phenyl group optionally substituted 1 to 5 times, wherein the substituents are independently selected from the group consisting of:

- a) -fluoro;
- b) -chloro;
- c) -bromo;

- d) -iodo;
- e) -cyano;
- f) -nitro; and
- g) -aryl.

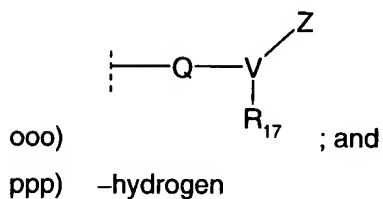
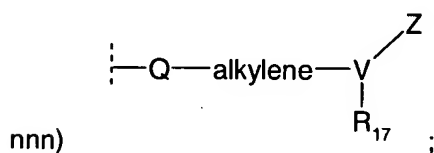
9. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein Ar₁ is a phenyl group substituted 1 to 5 times, wherein the substituents are selected from the group consisting of: -chloro ~~or~~ and -fluoro.

10. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein Ar₂ is a phenylene or naphthylene group optionally having 1 to 5 substituents, wherein the substituents are independently selected from the group consisting of:

- a) -fluoro;
- b) -chloro;
- c) -bromo;
- d) -iodo;
- e) -cyano;
- f) -nitro;
- g) -perfluoroalkyl;
- h) -Q-R₁₇;
- i) -alkyl;
- j) -aryl;
- k) -heteroaryl;
- l) -heterocyclyl;
- m) -cycloalkyl;
- n) -L₆-aryl;
- o) -L₆-arylene-aryl;
- p) -L₆-arylene-alkyl;
- q) -arylene-alkyl;
- r) -arylene-arylene-alkyl;
- s) -Q-alkyl;

- t) -Q-aryl;
- u) -Q-alkylene-aryl;
- v) -Q-arylene-alkyl;
- w) -Q-alkylene-arylene-aryl;
- x) -Q-arylene-arylene-aryl;
- y) -Q-alkylene-arylene-alkyl;
- z) -L₆-Q-alkylene-aryl;
- aa) -arylene-Q-alkyl;
- bb) -L₆-Q-aryl;
- cc) -L₆-Q-heteroaryl;
- dd) -L₆-Q-cycloalkyl;
- ee) -L₆-Q-heterocyclyl;
- ff) -L₆-Q-arylene-alkyl;
- gg) -L₆-Q-alkylene-arylene-alkyl;
- hh) -L₆-Q-alkyl;
- ii) -L₆-Q-alkylene-aryl-R₁₇;
- jj) -L₆-Q-alkylene-heteroaryl-R₁₇;
- kk) -arylene-Q-alkylene-R₁₇;
- ll) -heteroarylene-Q-alkylene-R₁₇;
- mm) -L₆-Q-aryl-R₁₇;
- nn) -L₆-Q-heteroarylene-R₁₇;
- oo) -L₆-Q-heteroaryl-R₁₇;
- pp) -L₆-Q-cycloalkyl-R₁₇;
- qq) -L₆-Q-heterocyclyl-R₁₇;
- rr) -L₆-Q-arylene-alkyl-R₁₇;
- ss) -L₆-Q-heteroarylene-alkyl-R₁₇;
- tt) -L₆-Q-alkylene-arylene-alkyl-R₁₇;
- uu) -L₆-Q-alkylene-heteroarylene-alkyl-R₁₇;
- vv) -L₆-Q-alkylene-cycloalkylene-alkyl-R₁₇;
- ww) -L₆-Q-alkylene-heterocyclylene-alkyl-R₁₇;
- xx) -L₆-Q-alkyl-R₁₇;
- yy) -L₆-Q-R₁₇;
- zz) -arylene-Q-R₁₇;

- aaa) -heteroarylene-Q-R₁₇;
- bbb) -heterocyclylene-Q-R₁₇;
- ccc) -Q-alkylene-R₁₇;
- ddd) -Q-arylene-R₁₇;
- eee) -Q-heteroarylene-R₁₇;
- fff) -Q-alkylene-arylene-R₁₇;
- ggg) -Q-alkylene-heteroarylene-R₁₇;
- hhh) -Q-heteroarylene-alkylene- R₁₇;
- iii) -Q-arylene-alkylene- R₁₇;
- jjj) -Q-cycloalkylene-alkylene- R₁₇;
- kkk) -Q-heterocyclylene-alkylene- R₁₇
- lll) -Q-alkylene-arylene-alkyl- R₁₇;
- mmm) -Q-alkylene-heteroarylene-alkyl- R₁₇;



wherein

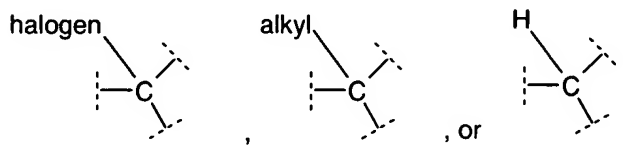
L₆ is a direct bond, -alkylene, -alkenylene, or -alkynylene;

Q is a direct bond, -CH₂-, -O-, -N(R₁₈)-, -C(O)-, -CON(R₁₈)-, -N(R₁₈)C(O)-, -N(R₁₈)CON(R₁₉)-, -N(R₁₈)C(O)O-, -OC(O)N(R₁₈)-, -N(R₁₈)SO₂-, -SO₂N(R₁₈)-, -C(O)-O-, -O-C(O)-, -S-, -S(O)-, -S(O₂)-, -N(R₁₈)SO₂N(R₁₉)-, -N=N-, or -N(R₁₈)-N(R₁₉)-;

wherein

R₁₈ and R₁₉ are independently selected from the group consisting of: -hydrogen, -alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, or and -alkylene-arylene-alkyl;

V is



Z is hydrogen, -alkylene-aryl, -alkyl, -aryl, -heteroaryl, -heterocyclyl, -cycloalkyl, -alkylene-heteroaryl, or -alkylene-cycloalkyl;

R₁₇ is -SO₃H, -P(O)(OH)₂, -P(O)(O-alkyl)(OH), -CO₂H, -CO₂-alkyl, an acid isostere, hydrogen, -alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, or -alkylene-arylene-alkyl.

11. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein Ar₂ is a phenyl group or naphthyl group optionally substituted 1 to 5 times, wherein the substituents are independently selected from the group consisting of:

- a) -fluoro;
- b) -chloro;
- c) -bromo;
- d) -iodo;
- e) -Q-R₁₇;
- f) -alkyl;
- g) -aryl;
- h) -arylene-alkyl;
- i) -Q-alkyl; and
- j) -arylene-Q-alkyl;

wherein

Q is -CH₂-, -O-, -C(O)-, or -C(O)-O-, and

R₁₇ is: -hydrogen, -alkyl, -aryl, -CO₂H, or an acid isostere.

12. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein Ar₂ is a phenyl group

substituted 1 to 5 times, wherein the substituents are independently selected from the group consisting of:

- a) -fluoro;
- b) -chloro;
- c) -bromo;
- d) -iodo;
- e) -Q-R₁₇;
- f) -alkyl;
- g) -phenyl;
- h) -phenylene-alkyl;
- i) -Q-alkyl; and
- j) -phenylene-Q-alkyl;

wherein

Q is: -CH₂-, -O-, -C(O)-, or -C(O)-O-, and

R₁₇ is: -hydrogen, -alkyl, -phenyl, or -CO₂H.

13. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein L₂ is: -CH₂-, -O-, alkylene, alkenylene, -O-alkylene-, -alkylene-O-, -N(R₂₀)-, -C(O)-, -CON(R₂₀)-, -N(R₂₀)C(O)-, -N(R₂₀)CON(R₂₁)-, -N(R₂₀)C(O)O-, -OC(O)N(R₂₀)-, -N(R₂₀)SO₂-, -SO₂N(R₂₀)-, -C(O)-O-, -O-C(O)-, -S-, -S(O)-, -S(O₂)-, -N(R₂₀)SO₂N(R₂₁)-, -N=N-, or -N(R₂₀)-N(R₂₁)- or a direct bond, wherein R₂₀ and R₂₁ independently selected from the group consisting of: -hydrogen, -alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, and -alkylene-arylene-alkyl.

14. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein L₂ is: -O-, -O-alkylene-, -alkylene-O-, or a direct bond.

15. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein L₂ is: -O-alkylene- or a direct bond.

16. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein T is an aryl group optionally having 1 to 5 substituents, wherein the substituents are independently selected from the group consisting of:

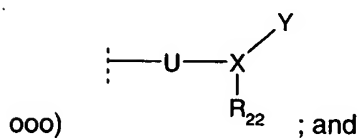
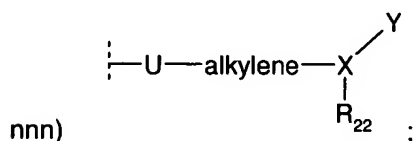
- a) -fluoro;
- b) -chloro;
- c) -bromo;
- d) -iodo;
- e) -cyano;
- f) -nitro;
- g) -perfluoroalkyl;
- h) -U-R₂₂;
- i) -alkyl;
- j) -aryl;
- k) -heteroaryl;
- l) -heterocyclyl;
- m) -cycloalkyl;
- n) -L₇-aryl;
- o) -L₇-arylene-aryl;
- p) -L₇-arylene-alkyl;
- q) -arylene-alkyl;
- r) -arylene-arylene-alkyl;
- s) -U-alkyl;
- t) -U-aryl;
- u) -U-alkylene-aryl;
- v) -U-arylene-alkyl;
- w) -U-alkylene-arylene-aryl;
- x) -U-arylene-arylene-aryl;
- y) -U-alkylene-arylene-alkyl;
- z) -L₇-U-alkylene-aryl;
- aa) -arylene-U-alkyl;
- bb) -L₇-U-aryl;
- cc) -L₇-U-heteroaryl;

- dd) -L₇-U-cycloalkyl;
- ee) -L₇-U-heterocyclyl;
- ff) -L₇-U-arylene-alkyl;
- gg) -L₇-U-alkylene-arylene-alkyl;
- hh) -L₇-U-alkyl;
- ii) -L₇-U-alkylene-aryl- R₂₂;
- jj) -L₇-U-alkylene-heteroaryl- R₂₂;
- kk) -arylene-U-alkylene- R₂₂;
- ll) -heteroarylene-U-alkylene- R₂₂;
- mm) -L₇-U-aryl- R₂₂;
- nn) -L₇-U-heteroarylene- R₂₂;
- oo) -L₇-U-heteroaryl- R₂₂;
- pp) -L₇-U-cycloalkyl- R₂₂;
- qq) -L₇-U-heterocyclyl- R₂₂;
- rr) -L₇-U-arylene-alkyl- R₂₂;
- ss) -L₇-U-heteroarylene-alkyl- R₂₂;
- tt) -L₇-U-alkylene-arylene-alkyl- R₂₂;
- uu) -L₇-U-alkylene-heteroarylene-alkyl- R₂₂;
- vv) -L₇-Q-alkylene-cycloalkylene-alkyl-R₂₂;
- ww) -L₇-Q-alkylene-heterocyclylene-alkyl-R₂₂;
- xx) -L₇-U-alkyl- R₂₂;
- yy) -L₇-U- R₂₂;
- zz) -arylene-U- R₂₂;
- aaa) -heteroarylene-U- R₂₂;
- bbb) -heterocyclylene-U- R₂₂;
- ccc) -U-alkylene- R₂₂;
- ddd) -U-arylene- R₂₂;
- eee) -U-heteroarylene- R₂₂;
- fff) -U-alkylene-arylene- R₂₂;
- ggg) -U-alkylene-heteroarylene- R₂₂;
- hhh) -U-heteroarylene-alkylene- R₂₂;
- iii) -U-arylene-alkylene- R₂₂;
- jjj) -U-cycloalkylene-alkylene- R₂₂;

kkk) -U-heterocyclylene-alkylene- R_{22} ;

lll) -U-alkylene-arylene-alkyl- R_{22} ;

mmm) -U-alkylene-heteroarylene-alkyl- R_{22} ;



ppp) -hydrogen;

wherein

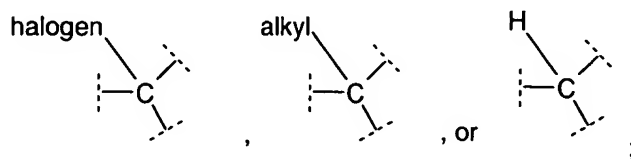
L_7 is a direct bond, -alkylene, -alkenylene, or -alkynylene;

U is a direct bond, $-\text{CH}_2-$, $-\text{O}-$, $-\text{N}(\text{R}_{23})-$, $-\text{C}(\text{O})-$, $-\text{CON}(\text{R}_{23})-$, $-\text{N}(\text{R}_{23})\text{C}(\text{O})-$, $-\text{N}(\text{R}_{23})\text{CON}(\text{R}_{24})-$, $-\text{N}(\text{R}_{23})\text{C}(\text{O})\text{O}-$, $-\text{OC}(\text{O})\text{N}(\text{R}_{23})-$, $-\text{N}(\text{R}_{23})\text{SO}_2-$, $-\text{SO}_2\text{N}(\text{R}_{23})-$, $-\text{C}(\text{O})-\text{O}-$, $-\text{O}-\text{C}(\text{O})-$, $-\text{S}-$, $-\text{S}(\text{O})-$, $-\text{S}(\text{O}_2)-$, $-\text{N}(\text{R}_{23})\text{SO}_2\text{N}(\text{R}_{24})-$, $-\text{N}=\text{N}-$, or $-\text{N}(\text{R}_{23})-\text{N}(\text{R}_{24})-$;

wherein

R_{23} and R_{24} are independently selected from the group consisting of: -hydrogen, -alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, and -alkylene-arylene-alkyl;

X is



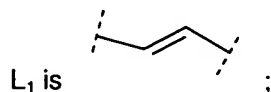
Y is hydrogen, -alkylene-aryl, -alkyl, -aryl, -heteroaryl, -heterocyclyl, -cycloalkyl, -alkylene-heteroaryl, or -alkylene-cycloalkyl;

R_{22} is $-\text{SO}_3\text{H}$, $-\text{P}(\text{O})(\text{OH})_2$, $-\text{P}(\text{O})(\text{O-alkyl})(\text{OH})$, $-\text{CO}_2\text{H}$, $-\text{CO}_2$ -alkyl, an acid isostere, -hydrogen, -alkyl, -aryl, -arylene-alkyl, -alkylene-aryl, or -alkylene-arylene-alkyl.

17. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein T is an aryl group substituted by -U-alkylene-R₂₂, wherein U is -O- or a direct bond, and R₂₂ is -CO₂H or an acid isostere.

18. (Currently Amended) The compound of Formula (I) according to claim 16 or a pharmaceutically acceptable salt thereof, wherein

a and b are equal to zero;



Ar₂ is a phenylene group optionally substituted 1 time with a group consisting of: -Q-alkyl, wherein Q is -O-;

L₂ is a direct bond, O-alkylene, or an -alkynylene; and

T is an aryl group substituted with at least one substituent selected from the group consisting of:

- a) -U-R₂₂;
- b) -U-alkylene-arylene-R₂₂;
- c) -U-alkylene-R₂₂;
- d) -U-arylene-R₂₂;
- e) -U-arylene-R₂₂ wherein the arylene is substituted with at least one of a halogen, methanesulfonylamino, or trifluoromethanesulfonylamino group [(.)] ;
- f) -U-arylene wherein the arylene is substituted with at least one trifluoromethanesulfonylamino group;
- g) -R_{22i}; and
- h) -halogen_i

wherein R₂₂ is -CO₂H or an acid isostere.

19. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein

a and b are equal to zero;

R₁ is hydrogen;

W is -N(R₂)-, wherein R₂ is alkyl; and

Ar₁ is aryl substituted 2 times wherein the substituent groups are -chloro.

20. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein W is -N(R₂)-, wherein R₂ is -L₃-D-alkylene-arylene-G, wherein L₃ is a direct bond or alkylene, D is a direct bond, or -O-, and G is -CN, -SO₃H, -P(O)(OH)₂, -P(O)(O-alkyl)(OH), -CO₂H, -CO₂-alkyl, or an acid isostere.

21. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein a and b are equal to 0, and T, L₂, Ar₂, and L₁ together form a group selected from a group consisting of:

(E)-2-(4-methoxyphenyl)vinyl, (E)-2-(3-methoxyphenyl)vinyl, (E)-2-(2-methoxyphenyl)vinyl, (E)-2-(3,4-dimethoxyphenyl)vinyl, (E)-2-(2,3,4-trimethoxyphenyl)vinyl, (E)-2-(4-ethoxyphenyl)vinyl, (E)-2-phenylvinyl, (E)-2-(4-fluorophenyl)vinyl, (E)-2-(4-chlorophenyl)vinyl, (E)-2-(4-bromophenyl)vinyl, (E)-2-(1,1'-biphenyl-4-yl)vinyl, (E)-2-(1-naphthyl)vinyl, (E)-2-(2-naphthyl)vinyl, 9H-fluoren-9-ylidenemethyl, (E)-2-(4'-methoxy-1,1'-biphenyl-4-yl)vinyl, (E)-2-(3'-methoxy-1,1'-biphenyl-4-yl)vinyl, (E)-2-(4-hydroxyphenyl)vinyl, 2-(4-methoxyphenyl)ethyl, (E)-2-(4'-carboxymethyloxy-1,1'-biphenyl-4-yl)vinyl, (E)-2-(4'-(3-methoxycarbonyl-1-propyloxy)-1,1'-biphenyl-4-yl)vinyl, (E)-2-(4'-(3-carboxy-1-propyloxy)-1,1'-biphenyl-4-yl)vinyl, (E)-2-(4'-phenoxy-1,1'-biphenyl-4-yl)vinyl, and (E)-2-(4'-benzyloxy-1,1'-biphenyl-4-yl)vinyl.

22. (Currently Amended) The compound of Formula (I) according to claim 1 or a pharmaceutically acceptable salt thereof, wherein Ar₁ is: 2,4-dichlorophenyl.

23. (Currently Amended) The compound of Formula (I) according to claim 1, where the compound of Formula (I) is:

4-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-3-fluorobiphenyl-4-yloxy)methyl)-benzoic acid;

4-(4-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-phenoxy)methyl)-benzoic acid;

4-[4'-(2-[4-(2,4-dichloro-phenyl)-1-[(1-naphthalen-1-yl-ethylcarbamoyl)-methyl]1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy]-butyric acid;

4-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-butyric acid;

5-[3-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-propyl]-1H-tetrazole;

[4-(3-(2-[4-(2,4-dichloro-phenyl)-1H-imidazol-2-yl]-(E)-vinyl)-4-methoxy-phenyl-ethynyl)-phenoxy]-acetic acid;

4-[3-(4-(2-[4-(2,4-dichloro-phenyl)-1H-imidazol-2-yl]-(E)-vinyl)-phenylethynyl)-phenoxy]-butyric acid;

5-[3-(4'-(2-[4-(2,4-dichloro-phenyl)-1-methyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-propyl]-1H-tetrazole;

5-(4'-(2-[4-(2,4-dichloro-phenyl)-1-methyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-pentanoic acid

2-bromo-4-(4'-(2-[4-(2,4-dichloro-phenyl)-1-methyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-benzoic acid;

4-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxymethyl)-benzoic acid;

4-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-benzoic acid;

2-bromo-4-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-benzoic acid;

4-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-3-methanesulfonylamino-benzoic acid;

4-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-3-trifluoromethanesulfonyl-amino-benzoic acid;

5-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-2-methanesulfonylamino-benzoic acid;

5-(4'-(2-[4-(2,4-dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-2-trifluoromethane-sulfonylamino-benzoic acid; or

4-(4'-(2-[4-(2,4-Dichloro-phenyl)-1-ethyl-1H-imidazol-2-yl]-(E)-vinyl)-biphenyl-4-yloxy)-butyric acid 2,2-dimethyl-propionyloxymethyl ester,
or a pharmaceutically acceptable salt thereof.

24. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable salt, solvate, or prodrug of a compound of Formula (I) according to a compound as claimed in claim 1.

25. (Currently Amended) The pharmaceutical composition of claim 24, wherein said ~~compound is applied to the skin~~ pharmaceutical composition is a topical formulation.

26. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in Claim 1 sufficient to inhibit protein tyrosine phosphatase.

27. (Original) The pharmaceutical composition of claim 26, in the form of an oral dosage or parenteral dosage unit.

28. (Original) The pharmaceutical composition of claim 26, wherein said compound is administered as a dose in a range from about 0.003 to 500 mg/kg of body weight per day.

29. (Original) The pharmaceutical composition of claim 26, wherein said compound is administered as a dose in a range from about 0.1 to 200 mg/kg of body weight per day.

30. (Original) The pharmaceutical composition of claim 26, wherein said compound is administered as a dose in a range from about 0.1 to 100 mg/kg of body weight per day.

31. (Currently Amended) The pharmaceutical composition of claim 26, further comprising one or more therapeutic agents selected from the group consisting of alkylating agents, antimetabolites, plant alkaloids, antibiotics, hormones, biologic response modifiers, analgesics, NSAIDs, DMARDs, glucocorticoids, sulfonylureas, biguanides, acarbose, PPAR agonists, DPP-IV inhibitors, GK activators, insulin, insulin mimetics, insulin secretagogues, insulin sensitizers, GLP-1, GLP-1 mimetics, cholinesterase inhibitors, antipsychotics, antidepressants, anticonvulsants, HMG CoA reductase inhibitors, cholestyramine, and fibrates. [[.]]

32. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat type I diabetes.

33. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat type II diabetes.

34. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat immune dysfunction.

35. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat AIDS.

36. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat autoimmune diseases.

37. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat glucose intolerance.

38. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat obesity.

39. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat cancer.

40. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat psoriasis.

41. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat allergic diseases.

42. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat infectious diseases.

43. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat inflammatory diseases.

44. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat diseases involving the modulated synthesis of growth hormone.

45. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat diseases involving the modulated synthesis of growth factors or cytokines which affect the production of growth hormone.

46. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a pharmacologically effective amount of the compound as claimed in claim 1, sufficient to treat Alzheimer's disease.

47. (Original) A method of inhibition protein tyrosine phosphatases which comprises administering to a subject in need thereof a pharmacologically effective amount of a compound as claimed in claim 1.

48. (Currently Amended) A method of prevention and/or treatment of PTPase mediated human diseases, treatment comprising alleviation of one or more symptoms resulting from that disorder, to an outright cure for that particular disorder or prevention of the onset of the disorder, the method comprising administration to a

human in need thereof a therapeutically effective amount of a compound of Formula (I) as claimed in claim 1.

49. (Original) The method of claim 47, further comprising administering to a subject in need thereof at least one adjuvant and/or additional therapeutic agent(s).

50. (Currently Amended) A method of treating PTPase mediated diseases, the method comprising administering to a subject in need thereof, a therapeutically effective amount of a compound of Formula (I) as claimed in claim 1, in combination with one or more therapeutic agents selected from the group consisting of alkylating agents, antimetabolites, plant alkaloids, antibiotics, hormones, biologic response modifiers, analgesics, NSAIDs, DMARDs, glucocorticoids, sulfonylureas, biguanides, acarbose, PPAR agonists, DPP-IV inhibitors, GK activators, insulin, insulin mimetics, insulin secretagogues, insulin sensitizers, GLP-1, GLP-1 mimetics, cholinesterase inhibitors, antipsychotics, antidepressants, anticonvulsants, HMG CoA reductase inhibitors, cholestyramine, and fibrates. [[.]]

51. (Currently Amended) A method for treating acute and/or chronic inflammation, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of Formula (I) as defined as claimed in claim 1.

52. (Currently Amended) A method for treating type I or type II diabetes, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of Formula (I) as defined as claimed in claim 1.

53. (Currently Amended) A method for treating immune dysfunction, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of Formula (I) as defined as claimed in claim 1.

54. (Currently Amended) A method for treating AIDS, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of Formula (I) as defined as claimed in claim 1.

55. (Currently Amended) A method for treating autoimmune disease, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of ~~Formula (I) as defined~~ as claimed in claim 1.

56. (Currently Amended) A method for treating glucose intolerance, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of ~~Formula (I) as defined~~ as claimed in claim 1.

57. (Currently Amended) A method for treating cancer, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of ~~Formula (I) as defined~~ as claimed in claim 1.

58. (Currently Amended) A method for treating psoriasis, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of ~~Formula (I) as defined~~ as claimed in claim 1.

59. (Currently Amended) A method for treating allergic diseases, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of ~~Formula (I) as defined~~ as claimed in claim 1.

60. (Currently Amended) A method for treating infectious disease, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of ~~Formula (I) as defined~~ as claimed in claim 1.

61. (Currently Amended) A method for treating diseases involving the modulated synthesis of growth hormone, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of ~~Formula (I) as defined~~ as claimed in claim 1.

62. (Currently Amended) A method for treating modulated synthesis of growth factors or cytokines which affect the production of growth hormone, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of ~~Formula (I) as defined~~ as claimed in claim 1.

63. (Currently Amended) A method for treating Alzheimer's disease, which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of Formula (I) ~~as defined~~ as claimed in claim 1.